1

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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
09/996,061	11/27/2001	Max Schaldach	7163-32	3174
21324	7590 03/30/2005		EXAM	INER
HAHN LOESER & PARKS, LLP One GOJO Plaza			THALER, MICHAEL H	
Suite 300	aza -		ART UNIT	PAPER NUMBER
AKRON, OF	I 44311-1076		3731	

DATE MAILED: 03/30/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

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	Application No.	Applicant(s)				
	09/996,061	SCHALDACH ET AL.				
Office Action Summary	Examiner	Art Unit				
	Michael Thaler	3731				
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet with	the correspondence address				
A SHORTENED STATUTORY PERIOD FOR REPI	I V IS SET TO EVOIDE 2 MC	NITH(S) EDOM				
THE MAILING DATE OF THIS COMMUNICATION  - Extensions of time may be available under the provisions of 37 CFR 1.  after SIX (6) MONTHS from the mailing date of this communication.  If the period for reply specified above is less than thirty (30) days, a replection of the period for reply is specified above, the maximum statutory period.  Failure to reply within the set or extended period for reply will, by statuth Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	. 136(a). In no event, however, may a rep ply within the statutory minimum of thirty d will apply and will expire SIX (6) MONTH te, cause the application to become ABA	oly be timely filed (30) days will be considered timely.  HS from the mailing date of this communication. NDONED (35 U.S.C. § 133).				
Status						
1) Responsive to communication(s) filed on 14 I	<u>March 2005</u> .					
,	·					
3) Since this application is in condition for allowa						
closed in accordance with the practice under	Ex parte Quayle, 1935 C.D.	11, 453 O.G. 213.				
Disposition of Claims						
4)⊠ Claim(s) <u>1-17 and 19-52</u> is/are pending in the	application.					
4a) Of the above claim(s) 7,9-13,19,20,35-40	4a) Of the above claim(s) 7,9-13,19,20,35-40 and 42-50 is/are withdrawn from consideration.					
5) Claim(s) is/are allowed.						
6)⊠ Claim(s) <u>1-6,8,14-17,21-34,41,51 and 52</u> is/a	re rejected.					
7) Claim(s) is/are objected to.						
8) Claim(s) are subject to restriction and/	or election requirement.	<i>C</i>				
Application Papers						
9) The specification is objected to by the Examin	er.					
10)☐ The drawing(s) filed on is/are: a)☐ ac	cepted or b) objected to by	y the Examiner.				
Applicant may not request that any objection to the	• • • • • • • • • • • • • • • • • • • •					
Replacement drawing sheet(s) including the corre	·	-				
11) The oath or declaration is objected to by the E	examiner. Note the attached	Office Action or form PTO-152.				
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreig a) All b) Some * c) None of:  1. Certified copies of the priority documer 2. Certified copies of the priority documer 3. Copies of the certified copies of the priority documer application from the International Burea * See the attached detailed Office action for a lis	nts have been received. nts have been received in Ap ority documents have been re au (PCT Rule 17.2(a)).	plication No eceived in this National Stage				
dec the attached detailed Office action for a lis	a c. the defined copies not re					
Attachment(s)						
1) Notice of References Cited (PTO-892)	4) Interview Su					
<ol> <li>Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08 Paper No(s)/Mail Date</li> </ol>		/Mail Date ormal Patent Application (PTO-152) -				

Application/Control Number: 09/996,061

Art Unit: 3731

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on March 1, 2005 has been entered.

Claims 7, 9-13, 19, 20, 35-40 and 42-50 are withdrawn from further consideration pursuant to 37 CFR 1.142(b), as being drawn to a nonelected invention and species, there being no allowable generic or linking claim. Applicant timely traversed the restriction (election) requirement in the reply filed on May 19, 2004.

Claims 1, 2, 5, 6, 25 and 30 are rejected under 35 U.S.C. 102(b) as anticipated by or, in the alternative, under 35 U.S.C. 103(a) as obvious over Turi (5,556,414). Turi discloses a stent 20 for a vessel (col. 1, lines 40-42) comprising a tubular body (figure 1) for expansion from a first condition to a second condition (col. 8, lines 1-5), the stent being configured such that a first part of the stent (e.g. the entire vein graft 26) is disposed inwardly (as well as outwardly) relative to a second part of the stent (e.g. member 22), (That

Art Unit: 3731

is, in the embodiments described in col. 5, lines 11-16 and 18-30, the vein graft 26 is disposed inwardly as well as outwardly relative to member 22.), and wherein in the second condition, at least a portion of the first part (e.g. portion 36 of vein graft 26) is not disposed inwardly relative to the second part 22 of the stent, wherein the tubular body includes at least a first wall portion (the wall of the entire composite prosthesis 20) comprising human or animal tissue (26) of adequate elasticity. Alternatively, it would have been obvious that the tissue 26 of the Turi stent 20 has adequate elasticity since it expands with the cylindrical member 22. As to claims 6 and 30, Turi discloses hardening agent (the adhesive described in col. 5, lines 49-52 which is a hardening agent since it hardens as it cures or dries).

Claims 4, 8, 22, 23, 27, 29, 32, 34 and 41 are rejected under 35 U.S.C. 103(a) as being unpatentable over Turi (5,556,414). As to claims 4, 22 and 23, Turi fails to disclose the tissue being genetically modified. However, it is old and well known in this art to genetically modify tissue in order to obtain favorable characteristics for it. It would have been obvious to genetically modify the Turi tissue so that it too would have this advantage. As to claims 8 and 41, Turi fails to disclose the hardening agent (adhesive) enclosed in

Art Unit: 3731

microcapsules. However, it is old and well known in this art to enclose adhesive in microcapsules in order to obtain the advantage of easily deploying the adhesive on the surface. It would have been obvious to enclose the Turi adhesive in microcapsules so that it too would have this advantage. The above well known in the art statements are taken to be admitted prior art because applicant failed to traverse the examiner's assertions (M.P.E.P. 2144.03).

Claims 3, 21, 24, 26, 28, 31 and 33 are rejected under 35 U.S.C. 103(a) as being unpatentable over Turi (5,556,414) in view of Atala (2003/0208279). Turi fails to disclose the tissue being cartilage. However, Atala teaches that tissue on a stent should be cartilage (paragraph [0041]) apparently in order to make the stent biocompatible (paragraph [0013]). It would have been obvious to make the Turi tissue cartilage so that it too would have this advantage.

Claims 14-17, 51 and 52 are rejected under 35 U.S.C. 103(a) as being unpatentable over Turi (5,556,414) in view of Berg et al. (5,680,873). As to claim 14, Turi discloses a catheter comprising a distal end region (the distal portion of the balloon catheter 41) and a holding device for holding the stent (the balloon on the balloon catheter 41). Turi fails to disclose a sheathing device provided with an application device

Art Unit: 3731

for applying a medium which is capable of flow to a surface of the stent. However, Berg et al. teach that a quide catheter 22 should be used with a balloon catheter in order to obtain the advantage of guiding the balloon catheter through vasculature as well as delivering fluids to the body (col. 1, It would have been obvious to include a quide lines 13-21). catheter with the Turi balloon catheter so that it too would have this advantage. Note that the Berg et al. guide catheter 22 (the claimed sheathing device) has an application device (the feed passage of guide catheter 22 through which dye passes as described in col. 7, lines 17-20) which is provided at the sheathing device for applying a medium which is capable of flow a surface of the stent. For example, after stent implantation, the balloon catheter could be removed from the quide catheter and die could be delivered through the quide catheter to the stent. As to claim 15, Berg et al. disclose an application opening (at the extreme distal end of guide catheter 22). As to claim 16, the Berg et al. sheathing device 22 has an anti-adhesion coating 40 while Turi discloses a adhesive in col. 5, lines 7-8 and 48-57. For this claim, the claimed stent may be considered to be only member 26 of Turi which includes a first part (the radially innermost portion of member 26) and a second part (the radially outermost portion of

Application/Control Number: 09/996,061

Art Unit: 3731

member 26). Note that the layer of adhesive on member 26 is on

the surface of member 26 facing radially outwardly (i.e. toward

the sheathing device as claimed).

Applicant's arguments filed March 1, 2005 have been fully

considered but they are not persuasive for the reasons set forth

above.

Any inquiry concerning this communication or earlier

communications from the examiner should be directed to Michael

Thaler whose telephone number is (571)272-4704. The examiner

can normally be reached Monday to Friday.

Ιf attempts to reach the examiner by telephone are

unsuccessful, the examiner's supervisor, Anhtuan T. Nguyen can

be reached on (571)272-4963. The fax phone number for the

organization where this application or proceeding is assigned is

(703)872 - 9306.

mht

3/22/05

PRIMARY EXAMINER

ART UNIT 3731

Page 6